FDA 510(k) Premarket Notification

K132066 INFLAMMA DRY

CLEARED

Nov 21, 2013
141 days to decision

K132066 · Product code: **PFQ** · Chemistry
Source: https://www.510kdatabase.net/k132066/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mmp-9 Test System (PFQ)
Date received	Jul 3, 2013
Decision date	Nov 21, 2013
Days to decision	141 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Rapid Pathogen Screening, Inc.**
Location	Sarasota, FL, US
Contact	Douglas Bueschel
510(k) history	2 submissions · 2 cleared · 2011-2013